AMENDMENTS TO THE CLAIMS

- 1. 10. (Canceled)
- 11. (Currently amended) A process for the culture of culturing mouse pluripotent stem cells, which comprises culturing the mouse pluripotent stem cells in a medium comprising leukemia inhibitory factor (LIF), an antioxidant and an inhibitor of adenylate cyclase activity under a condition such that adenylate cyclase activity is inhibited, said process allowing the mouse pluripotent stem cells to proliferate or establish while maintaining the cells in an undifferentiated state.
 - 12. (Canceled).
- 13. (Previously presented) The process according to claim 11, wherein the culture process is performed using a minimal culture medium.
- 14. (Previously presented) The process according to claim 11, wherein the pluripotent stem cells are ES cells.
 - 15. (Canceled)
 - 16. (Canceled)
- 17. (Currently amended) A process for the preparation of a clonal population of undifferentiated <u>mouse</u> pluripotent stem cells, which comprises culturing the undifferentiated <u>mouse</u> pluripotent stem cells <u>in a medium comprising leukemia inhibitory factor (LIF)</u>, an <u>antioxidant and an inhibitor of adenylate cyclase activity under a condition such that adenylate cyclase activity is inhibited</u>.
- 18. (Currently amended) A process for the preparation of a clonal population of undifferentiated <u>mouse</u> pluripotent stem cells, which comprises isolating undifferentiated pluripotent stem cells from a living body of a mouse, and culturing the undifferentiated <u>mouse</u> pluripotent stem cells in a medium comprising leukemia inhibitory factor (LIF), an antioxidant

and an inhibitor of adenylate cyclase activityunder a condition such that adenylate cyclase activity is inhibited.

- 19. (Canceled).
- 20. (Previously presented) The process according to claim 17, wherein the culture process is performed using a minimal culture medium.
- 21. (Previously presented) The process according to claim 17, wherein one pluripotent stem cell is cultured to provide a clonal population of the cells.
- 22. (Previously presented) The process according to claim 17, wherein pluripotent stem cells are cultured in a medium free of feeder cells or free of serum or free of both, to provide a clonal population of the cells, in which the pluripotent stem cells are seeded at a lower density than that which allows adjacent pluripotent stem cells to interact with each other, so as to induce the proliferation of undifferentiated pluripotent stem cells.
- 23. (Previously presented) The process according to claim 17, wherein one pluripotent stem cell is cultured in a medium free of feeder cells or free of serum or free of both, to provide a clonal population of the cells.
- 24. (Previously presented) The process according to claim 17, wherein the pluripotent stem cells are ES cells.
 - 25. (Canceled)
 - 26. (Canceled)
 - 27. -29. (Canceled)
- 30. (Currently amended) The process of claim 12 claim 11, wherein the inhibitor of adenylate cyclase activity is selected from the group consisting of SQ22536 (9-(tetrahydro-2-furanyl)adenine), 2',5'-dideoxyadenosine, 9-cyclopentyladenine, 2',5'-dideoxyadenosine 3'-

diphosphate, 2',5'-dideoxyadenosine 3'-monophosphate, and MDL-12,330A (cis-N-(2-phenylcyclopentyl)azacyclotridec-1-en-2-amine).

31. (Canceled)

32. (Currently amended) The process of-claim 12claim 11, wherein the inhibitor of adenylate cyclase activity is selected from the group consisting of adrenocorticotropic hormone (ACTH), brain natriuretic peptide (BNP), pituitary adenylate cyclase activating polypeptide (PACAP), and a peptide having a physiological activity substantially similar to them.

33. (Canceled)

- 34. (Previously presented) The process according to claim 11, wherein the medium is free of feeder cells or of serum or free of both.
- 35. (Previously presented) The process according to claim 11, wherein the medium further comprises a differentiation inhibitory factor, a serum replacement and an antioxidant.
- 36. (Previously presented) The process according to claim 17, wherein the medium further comprises a differentiation inhibitory factor, a serum replacement and an antioxidant.
- 37. (Currently amended) The process of <u>elaim 12 claim 11</u>, wherein the medium comprises leukemia inhibitory factor, 2-mercaptoethanol, KSR and adrenocorticotrophic hormone.

38. (Canceled)

39. (Currently amended) The process of <u>elaim 12 claim 11</u>, wherein the medium comprises leukemia inhibitory factor, 2-mercaptoethanol, KSR and SQ22536.

40. (Canceled)